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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
08/479,038	06/07/1995	William N. Drohan	1327.0440006	7774	
75	7590 07/13/2005			EXAMINER	
STERNE KESSLER GOLDSTEIN AND FOX, P.L.L.C.			MARSCHEL, ARDIN H		
SUITE 600	KK AVENUE N W		ART UNIT	PAPER NUMBER	
WASHINGTO	N, DC 20005-3934		1631		

DATE MAILED: 07/13/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)	
,	08/479,038	DROHAN ET AL.	
Office Action Summary	Examiner	Art Unit	
	Ardin Marschel	1631	
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the c	orrespondence address	
A SHORTENED STATUTORY PERIOD FOR REPL' THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period of the period for reply within the set or extended period for reply will, by statute the period of the	36(a). In no event, however, may a reply be tin y within the statutory minimum of thirty (30) day will apply and will expire SIX (6) MONTHS from t, cause the application to become ABANDONE	nely filed s will be considered timely. the mailing date of this communication D (35 U.S.C. § 133).	I.
Status		·	
1)⊠ Responsive to communication(s) filed on <u>02 M</u>	larch 2005.	•	
	action is non-final.		
3) Since this application is in condition for allowar	nce except for formal matters, pro	secution as to the merits is	i
closed in accordance with the practice under E	Ex parte Quayle, 1935 C.D. 11, 45	53 O.G. 213.	
Disposition of Claims	•		
4)⊠ Claim(s) <u>12,13,17-20,24-32 and 34-38</u> is/are p	ending in the application.	·	
4a) Of the above claim(s) is/are withdraw	• • • • • • • • • • • • • • • • • • • •		
5) Claim(s) is/are allowed.			
6) Claim(s) 12,13,17-20,24-32 and 34-38 is/are re	ejected.		
7) Claim(s) is/are objected to.			
8) Claim(s) are subject to restriction and/o	r election requirement.		
Application Papers			,
9) The specification is objected to by the Examine	r		
10)⊠ The drawing(s) filed on <u>11/1/04</u> is/are: a)⊠ acc		Examiner.	
Applicant may not request that any objection to the			
Replacement drawing sheet(s) including the correct			).
11) The oath or declaration is objected to by the Ex			,
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign	priority under 35 U.S.C. § 119(a)	-(d) or (f).	
a) All b) Some * c) None of:	- h h		
1. Certified copies of the priority documents		N-	
2. Certified copies of the priority documents	• •		
<ol> <li>Copies of the certified copies of the prior application from the International Bureau</li> </ol>	•	d in this National Stage	
* See the attached detailed Office action for a list		d	
occ the attached detailed office action for a list	or the certified copies hot receive	u.	
and a city			
Attachment(s)	A) 🗖 1-4 🛕	(DTO 440)	
Notice of References Cited (PTO-892)     Notice of Draftsperson's Patent Drawing Review (PTO-948)	4) LInterview Summary Paper No(s)/Mail Da		
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date		atent Application (PTO-152)	
J.S. Patent and Trademark Office PTOL-326 (Rev. 1-04)  Office Ac	tion Summary	Part of Paper No./Mail Date 7100	5

## **DETAILED ACTION**

Applicants' arguments, filed 3/2/05, have been fully considered but they are not deemed to be persuasive. Rejections and/or objections not reiterated from previous office actions are hereby withdrawn. The following rejections and/or objections are either reiterated or newly applied. They constitute the complete set presently being applied to the instant application.

## **NEW MATTER**

Claims 12, 13, 17-20, 24-32, and 34-38 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention.

This rejection is reiterated and maintained from the previous office action, mailed 7/2/04, and as necessitated by amendment due to the newly added claim 38. The NEW MATTER determination regarding the phrase "said effective amount of said supplement is greater than the amount which is soluble in said fibrin matrix" is maintained in that there NEW MATTER regarding generic supplement is maintained as well as the fibrin matrix content being based on "insolubility" and not on what is soluble as claimed as well as supplements characterized via molecular weight compared to ethanol.

Applicants point to page 22, lines 4-16, for support. Consideration of said page 22 citation reveals that only the inclusion of compounds such as free TET etc. confers extended longevity which can be exploited to increase duration of a drug's release. This

Art Unit: 1631

fails to provide written basis for any solubility or insolubility practice whatsoever, nor such solubility or insolubility in a fibrin matrix. This argument therefore is non-persuasive as clearly not providing written basis for the above NEW MATTER phrase. Applicants go on to point to page 104, lines 27-29, for written basis. Consideration of said page 104 citation reveals that lower solubility of antibiotics results in release over longer periods of time than highly soluble preparations. No written basis is present therein for solubility greater than in a fibrin matrix which is a requirement of the above cited NEW MATTER phrase and thus is also non-persuasive.

NEW MATTER has been amended into the claims via the citation of "said composition" being "substantially free of protease inhibitors" (claim 12, 34, and 36). This phrase has not been found as filed nor pointed to for support by applicants. It is acknowledged that supplements which do not have "added" protease inhibitor(s) is disclosed, such as in the original specification on page 41, lines 6-10, but that the supplement itself being substantially free of protease inhibitors has not been disclosed as filed.

## **PRIOR ART**

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein

Art Unit: 1631

were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 17, 18, 25, 29-32, and 35-38 are rejected under 35 U.S.C. 103(a) as being unpatentable over Marx (P/N 5,607,694); taken in view of Popescu et al. (P/N 4,708,861).

This rejection is maintained and reiterated from the previous office action, mailed 7/2/04. Applicants argue that priority should be granted to an earlier filed parent. This is not persuasive because the above NEW MATTER rejections prevent such priority granting as written basis has not been found to support such priority granting.

Applicants further argue regarding Popescu et al. In response, firstly the rejection is based on the combination of references and not on Popescu et al. alone. Secondly, in response, the liposome-containing supplements reasonably will vastly slow release thereof as being containment entities. Thus, release will clearly be slowed due to said containment which also prevents liquid in the area of the liposomes from entering to solubilize drugs therein and thus will result in a sustained period of release that is greater than that obtained due to solubility in the fibrin matrix which is outside of the liposomes and not in direct contact with any liposome contained drug. Thus, this rejection is maintained.

No claim is allowed.

Art Unit: 1631

Applicants' amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicants are reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Since the fee set forth in 37 CFR 1.17(r) for a first submission subsequent to a final rejection has been previously paid, applicant, under 37 CFR 1.129(a), is entitled to have a second submission entered and considered on the merits if, prior to abandonment, the second submission and the fee set forth in 37 CFR 1.17(r) are filed prior to the filing of an appeal brief under 37 CFR 1.192. Upon the timely filing of a second submission and the appropriate fee for a small entity under 37 CFR 1.17(r), the finality of the previous Office action will be withdrawn. If a notice of appeal and the appeal fee set forth in 37 CFR 41.20(b) were filed prior to or with the payment of the fee set forth in 37 CFR 1.17(r), the payment of the fee set forth in 37 CFR 1.17(r) by applicant will be construed as a request to dismiss the appeal and to continue

Art Unit: 1631

prosecution under 37 CFR 1.129(a). In view of 35 U.S.C. 132, no amendment considered as a result of payment of the fee set forth in 37 CFR 1.17(r) may introduce new matter into the disclosure of the application.

Papers related to this application may be submitted to Technical Center 1600 by facsimile transmission. Papers should be faxed to Technical Center 1600 via the Central PTO Fax Center. The faxing of such papers must conform with the notices published in the Official Gazette, 1096 OG 30 (November 15, 1988), 1156 OG 61 (November 16, 1993), and 1157 OG 94 (December 28, 1993)(See 37 CFR § 1.6(d)). The Central PTO Fax Center number is (571) 273-8300.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ardin Marschel, Ph.D., AU 1631 Supervisory Patent Examiner, whose telephone number is (571) 272-0718. The examiner can normally be reached on Monday-Friday from 8 A.M. to 4 P.M.

Any inquiry of a general nature or relating to the status of this application should be directed to Legal Instrument Examiner, Tina Plunkett, whose telephone number is (571) 272-0549.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

July 11, 2005

SUPERVISORY PATENT EXAMINER